

(FILE 'HOME' ENTERED AT 16:13:54 ON 15 MAR 2002)

FILE 'BIOSIS, EMBASE, CAPLUS' ENTERED AT 16:14:04 ON 15 MAR 2002

L1 13871 S TERATOMA
L2 51 S L1 AND (EMBRYONIC STEM CELL# OR EMBRYONIC DISK CELL# OR
EMBYR
L3 38 DUP REM L2 (13 DUPLICATES REMOVED)
L4 5 S L1 AND IMMUNOCOMPETENT
L5 3 DUP REM L4 (2 DUPLICATES REMOVED)
L6 0 S L1 AND BALBC
L7 27 S L1 AND BALB C
L8 19 DUP REM L7 (8 DUPLICATES REMOVED)

=> d 18 17 au ti so ab

L8 ANSWER 17 OF 19 EMBASE COPYRIGHT 2002 ELSEVIER SCI. B.V.

AU Siegler E.L.; Tick N.; Teresky A.K.; et al.

TI Teratocarcinoma transplantation rejection loci: An H-2-linked tumor
rejection locus.

SO Immunogenetics, (1979) 9/3 (207-220).

CODEN: IMNGBK

AB Embryoid bodies (ascites tumor) from a 129/Sv transplantable teratocarcinoma produce tumors (100%) in syngenic 129/Sv mice but fail to form tumors (3-6%) in **BALB/c** mice, C3H/He mice and C57BL/6 mice, in spite of the fact that the malignant stem cells of this tumor do not express detectable H-2 antigens. The available evidence indicates that this allogeneic tumor restriction has an immunological basis; 100% of the F1 hybrid mice between 129/Sv and the three other inbred mouse strains accept the 129/Sv teratocarcinoma. The backcross and F2 mice segregate the **BALB/c**, C3H/He and C57BL/6 tumor transplantation rejection loci in a manner that indicates that each of these inbred strains of mice contain one to two major transplantation rejection loci. A linkage analysis in the **BALB/c** and C3H/He backcross and F2 generations indicates that these mice have a teratocarcinoma transplantation rejection locus on chromosome 17, about eight to nine recombination units from the H-2 complex. An F1 complementation analysis between allogeneic mice that each reject teratocarcinomas tumors (BALB/cxC57BL/6 and C3H/HexC57BL/6), indicates that the C57BL/6 mice have the 129/Sv tumor-accepting (sensitive) allele at the H-2-linked locus but reject teratocarcinomas because of antigenic differences at a second locus. While these major teratocarcinoma transplantation rejection loci determine the acceptance or rejection of a tumor by a mouse injected with high doses of tumor tissue (750. μ g of tumor protein) evidence is presented for a number of minor genetic factors

that can (1) affect the efficiency of tumor rejection and (2) cause complete tumor rejection at lower tumor doses (7.5-75. μ g of tumor protein).

- L8 ANSWER 1 OF 19 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
1
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Okamura, Yasuyuki; Matsumoto, Kiyoshi
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cell line (TES-1).
- SO Anticancer Research, (May June, 1999) Vol. 19, No. 3A, pp. 1933-1940.
ISSN: 0250-7005.
- L8 ANSWER 2 OF 19 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
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- TI Genetic regulation of traits essential for spontaneous ovarian
teratocarcinogenesis in strain LT/Sv mice: Aberrant meiotic cell cycle,
oocyte activation, and parthenogenetic development.
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- L8 ANSWER 3 OF 19 BIOSIS. COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
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Ken-Ichi; Watanabe, Yoshihisa; Nagata, Emi; Nakata, Minoru
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mouse embryo in vitro.
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- AU VAN BERLO R J; DÉ JONG B; OOSTERHUIS J W; DIJKHUIZEN T; BUIST J; DAM A
- TI CYTOGENETIC ANALYSIS OF MURINE EMBRYO-DERIVED TUMORS.
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- L8 ANSWER 5 OF 19 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
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- TI YOLK-SAC CARCINOMA DEVELOPS SPONTANEOUSLY AS A LATE OCCURRENCE IN
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- L8 ANSWER 6 OF 19 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
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- TI CHARACTERIZATION OF TRANSFORMING GROWTH FACTORS PRODUCED BY THE
INSULIN-INDEPENDENT **TERATOMA**-DERIVED CELL LINE 1246-3A.
- SO J CELL PHYSIOL, (1989) 140 (2), 254-263.
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- L8 ANSWER 7 OF 19 EMBASE COPYRIGHT 2002 ELSEVIER SCI. B.V.
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- TI Alternative modes of c-myc regulation in growth factor-stimulated and
differentiating cells.

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LAFERTE

S; KRANTZ M J

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trophoblast cell clones.

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